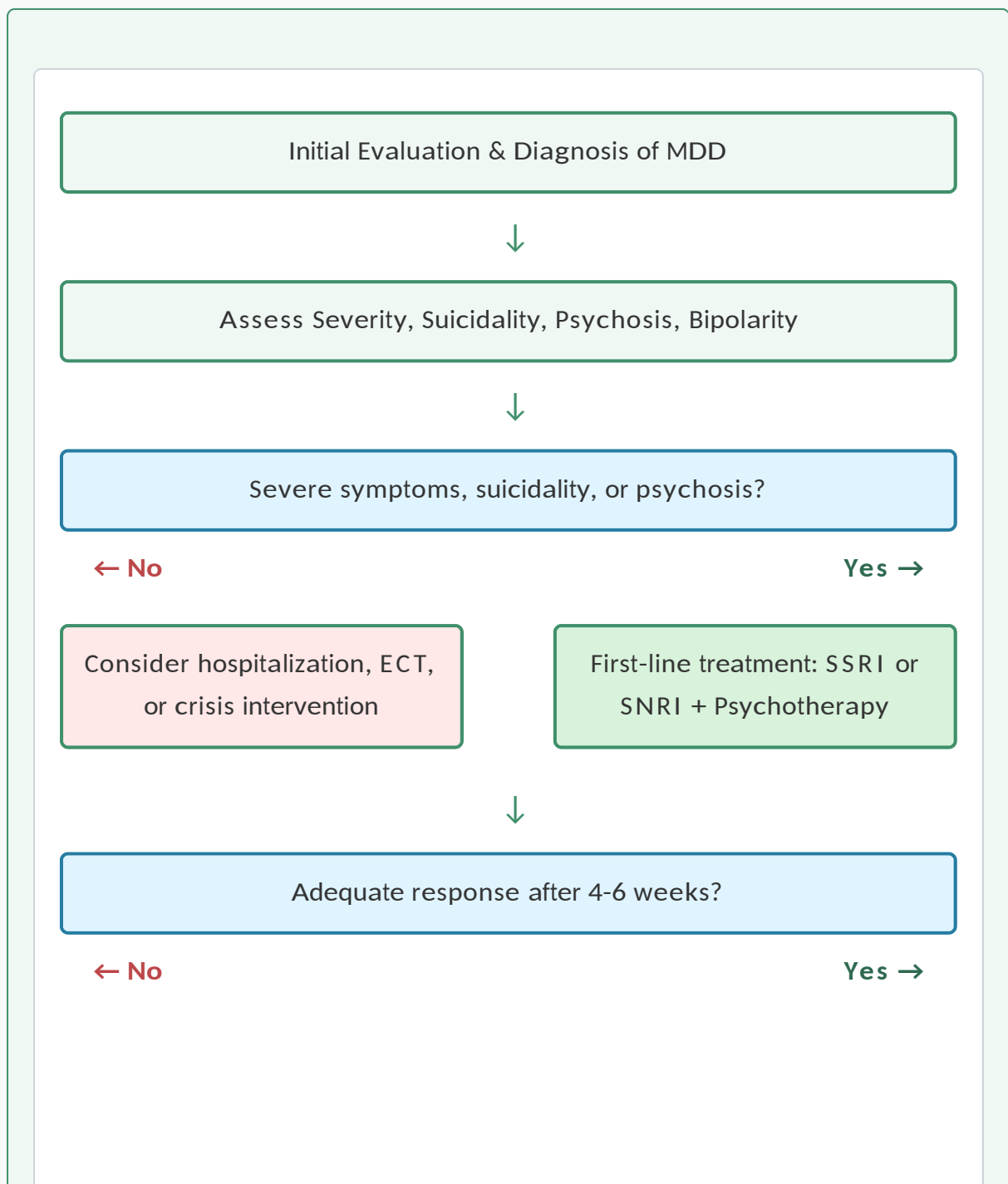


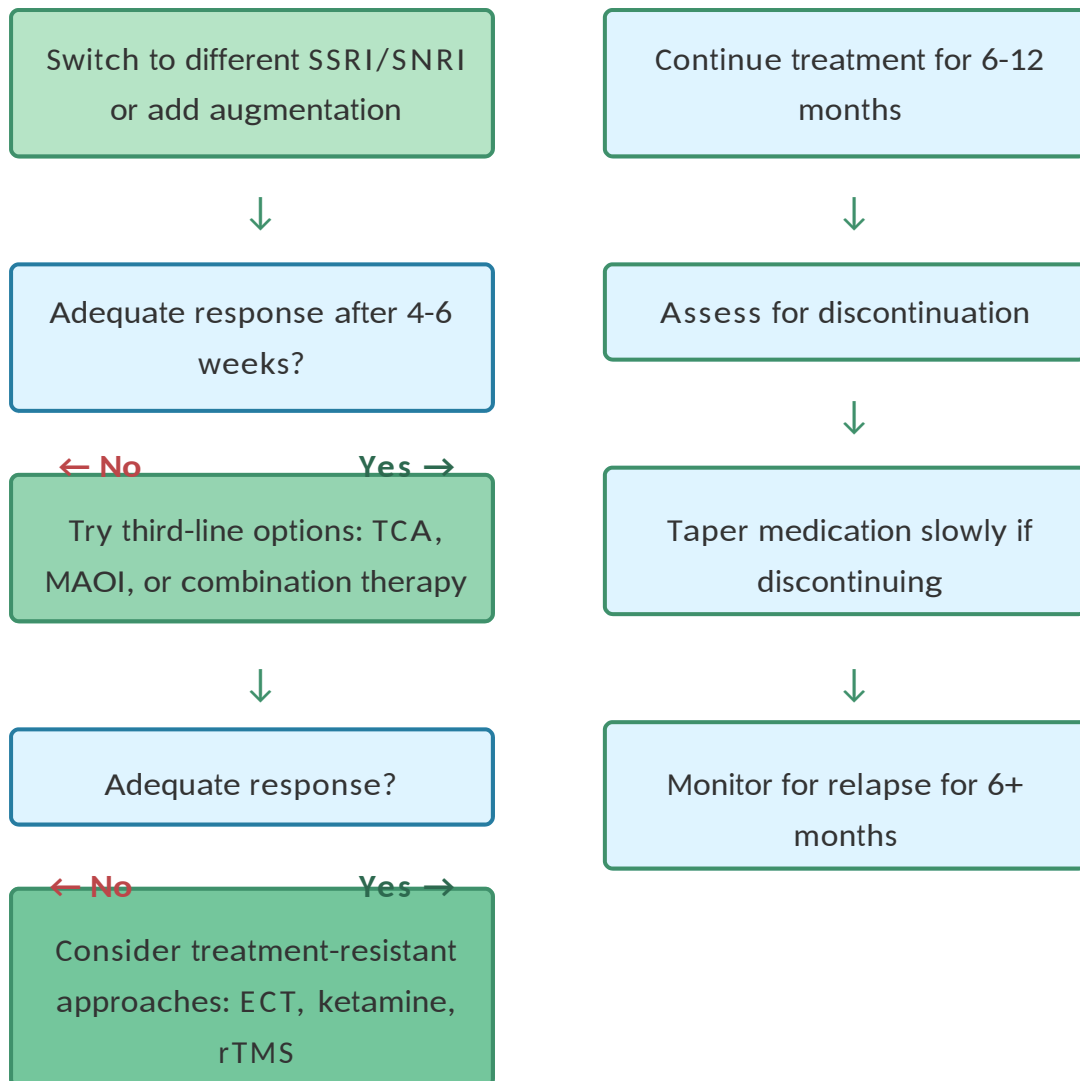


# Evidence-Based Treatment Algorithms for Psychiatric Disorders



## Major Depressive Disorder (MDD) Treatment Algorithm





### First-Line Treatment Options

**SSRIs:** Sertraline, Escitalopram, Fluoxetine, Citalopram

**SNRIs:** Venlafaxine, Duloxetine

**Psychotherapy:** Cognitive Behavioral Therapy (CBT), Interpersonal Therapy (IPT)

### Second-Line Treatment Options

Switch to different SSRI/SNRI

### Augmentation strategies:

- Add bupropion
- Add mirtazapine
- Add second-generation antipsychotic (aripiprazole, quetiapine, brexpiprazole)
- Add lithium
- Add thyroid hormone (T3)

### Third-Line Treatment Options

**Tricyclic Antidepressants (TCAs):** Nortriptyline, Amitriptyline

**Monoamine Oxidase Inhibitors (MAOIs):** Phenelzine, Tranylcypromine

**Combination therapy:** SSRI + Mirtazapine, SSRI + TCA

### Treatment-Resistant Depression Options

**Electroconvulsive Therapy (ECT)**

**Ketamine/Esketamine**

**Repetitive Transcranial Magnetic Stimulation (rTMS)**

**Vagus Nerve Stimulation (VNS)**

**Deep Brain Stimulation (DBS)**

### Monitoring Requirements

**Baseline:** Complete blood count, comprehensive metabolic panel, thyroid function tests

**Follow-up:** Assess response at 2, 4, 6, 8, and 12 weeks

**Ongoing:** Monitor for suicidality, especially in adolescents and young adults

**Specific monitoring:**

- TCAs: ECG, blood levels

- Lithium: Lithium levels, renal and thyroid function
- Antipsychotics: Metabolic parameters, movement disorders

### Medication Dosing Guidelines

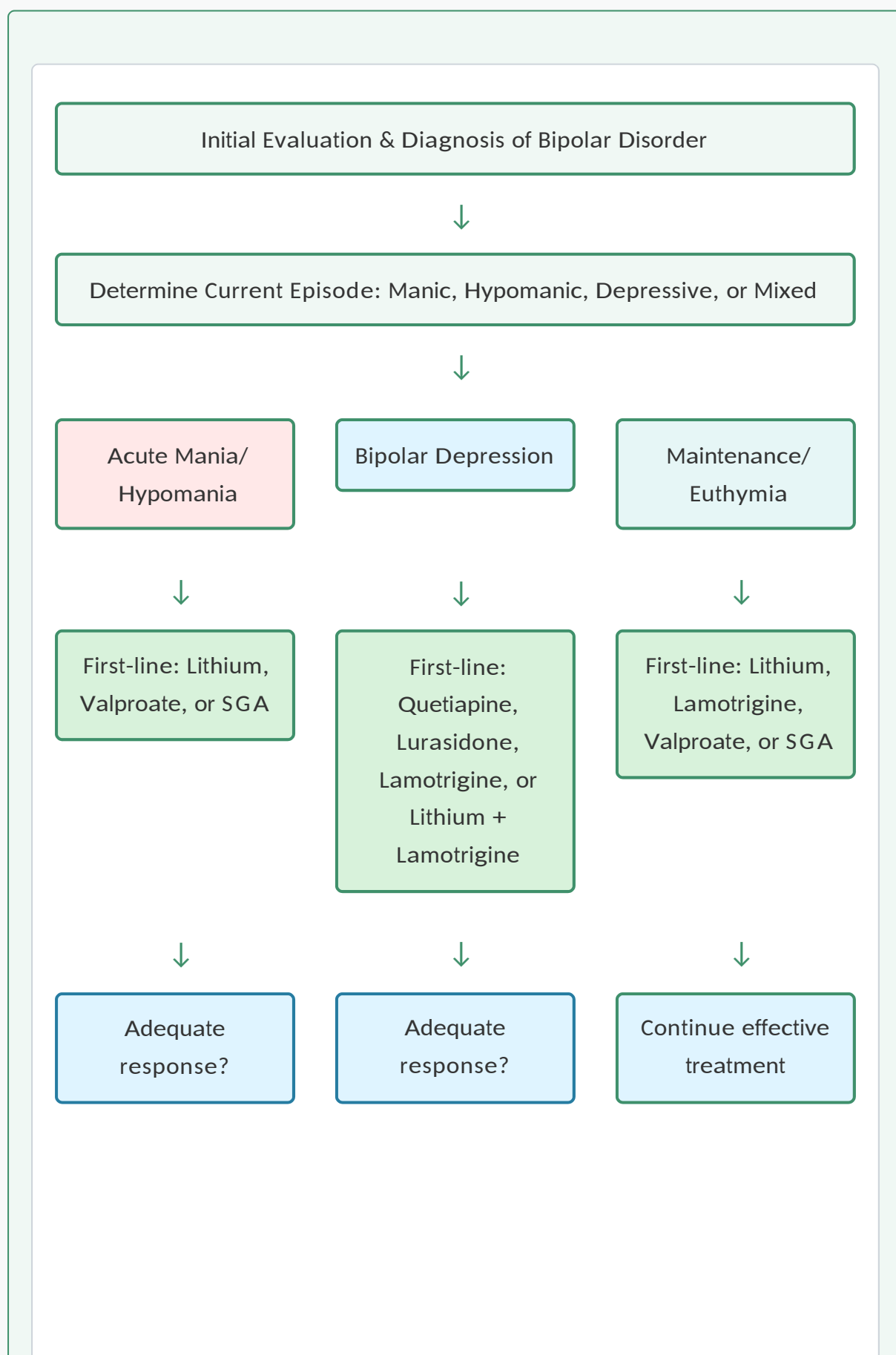
Medication	Starting Dose	Therapeutic Dose Range	Maximum Dose
Sertraline	25-50 mg/day	50-200 mg/day	200 mg/day
Escitalopram	5-10 mg/day	10-20 mg/day	20 mg/day
Fluoxetine	10-20 mg/day	20-60 mg/day	80 mg/day
Venlafaxine XR	37.5-75 mg/day	150-225 mg/day	375 mg/day
Duloxetine	30 mg/day	60-120 mg/day	120 mg/day
Bupropion XL	150 mg/day	300 mg/day	450 mg/day
Mirtazapine	15 mg/day	30-45 mg/day	45 mg/day
Aripiprazole (augmentation)	2-5 mg/day	5-15 mg/day	15 mg/day

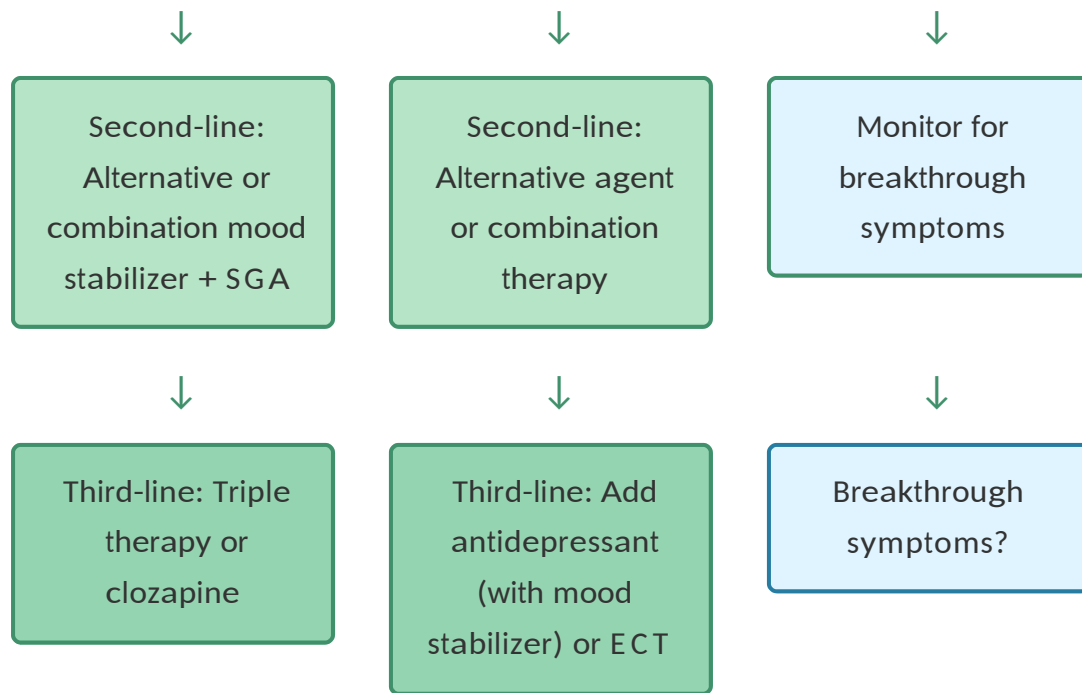
**Warning: Always assess for suicidality before and during treatment. Monitor closely during the first few weeks of treatment and with dose changes. Use caution when prescribing TCAs and MAOIs due to toxicity in overdose and dietary/drug interactions.**

**Clinical Pearl:** Antidepressant response may take 4-6 weeks for full effect. Partial response at 2-4 weeks often predicts eventual response. Consider psychotherapy in combination with medication for better outcomes, especially for mild to moderate depression.



## Bipolar Disorder Treatment Algorithm





### Acute Mania/Hypomania Treatment

#### First-line options:

- Lithium
- Valproate
- Second-generation antipsychotics (SGAs): Olanzapine, Risperidone, Quetiapine, Aripiprazole, Ziprasidone, Asenapine

#### Second-line options:

- Carbamazepine
- Lithium + SGA
- Valproate + SGA
- Lithium + Valproate

#### Third-line options:

- Triple therapy (Lithium + Valproate + SGA)
- Clozapine
- Electroconvulsive therapy (ECT)

## Bipolar Depression Treatment

### First-line options:

- Quetiapine
- Lurasidone (with lithium or valproate)
- Lamotrigine
- Lithium
- Olanzapine-fluoxetine combination

### Second-line options:

- Valproate
- Lithium + Lamotrigine
- Adjunctive psychotherapy (CBT, family-focused, interpersonal)

### Third-line options:

- Cariprazine
- Olanzapine
- Lithium or valproate + SSRI/bupropion (with careful monitoring)
- Electroconvulsive therapy (ECT)

## Maintenance Treatment

### First-line options:

- Lithium
- Lamotrigine (especially for depression prevention)
- Valproate
- Quetiapine
- Aripiprazole

### Second-line options:

- Olanzapine
- Risperidone LAI

- Ziprasidone
- Carbamazepine

**Warning: Antidepressant monotherapy is contraindicated in bipolar disorder due to risk of mood switch. Always use antidepressants with mood stabilizers if needed. Discontinue antidepressants after acute bipolar depression resolves.**

### Medication Dosing Guidelines

Medication	Starting Dose	Therapeutic Dose Range	Therapeutic Blood Level
Lithium	300 mg BID or TID	900-1800 mg/day	0.6-1.2 mEq/L (maintenance) 0.8-1.2 mEq/L (acute mania)
Valproate	250-500 mg BID	750-2500 mg/day	50-125 µg/mL
Lamotrigine	25 mg/day (titrate slowly)	100-400 mg/day	Not established
Carbamazepine	100-200 mg BID	600-1600 mg/day	4-12 µg/mL
Quetiapine	50 mg BID	300-800 mg/day	Not established
Olanzapine	5-10 mg/day	5-20 mg/day	Not established
Risperidone	1-2 mg/day	2-6 mg/day	Not established
Aripiprazole	5-15 mg/day	15-30 mg/day	Not established

### Monitoring Requirements

#### Lithium:

- Baseline: CBC, electrolytes, BUN/Cr, TSH, ECG (>40 years)



- Lithium levels: 5-7 days after initiation/dose change, then every 3-6 months
- Renal and thyroid function: Every 6-12 months

#### Valproate:

- Baseline: LFTs, CBC with platelets, pregnancy test
- LFTs and CBC: After 1 month, then every 3-6 months
- Valproate levels: As clinically indicated

#### Carbamazepine:

- Baseline: CBC, LFTs, electrolytes, pregnancy test
- CBC and LFTs: Monthly for first 3 months, then every 6-12 months
- Carbamazepine levels: As clinically indicated

#### Antipsychotics:

- Baseline: Weight, BMI, waist circumference, BP, fasting glucose, lipid panel
- Weight: Monthly for 3 months, then quarterly
- Metabolic parameters: At 3 months, then annually
- AIMS for EPS: Every 6 months

**Clinical Pearl:** Lithium remains the gold standard for suicide prevention in bipolar disorder. Consider predominant polarity (manic vs. depressive) when selecting maintenance treatment. Lamotrigine is more effective for preventing depression, while lithium and antipsychotics are more effective for preventing mania.



## Schizophrenia Treatment Algorithm

Initial Evaluation & Diagnosis of Schizophrenia



Assess Symptom Severity, Suicidality, Comorbidities



Acute psychosis requiring immediate intervention?

← No

Yes →

Consider hospitalization, IM antipsychotic

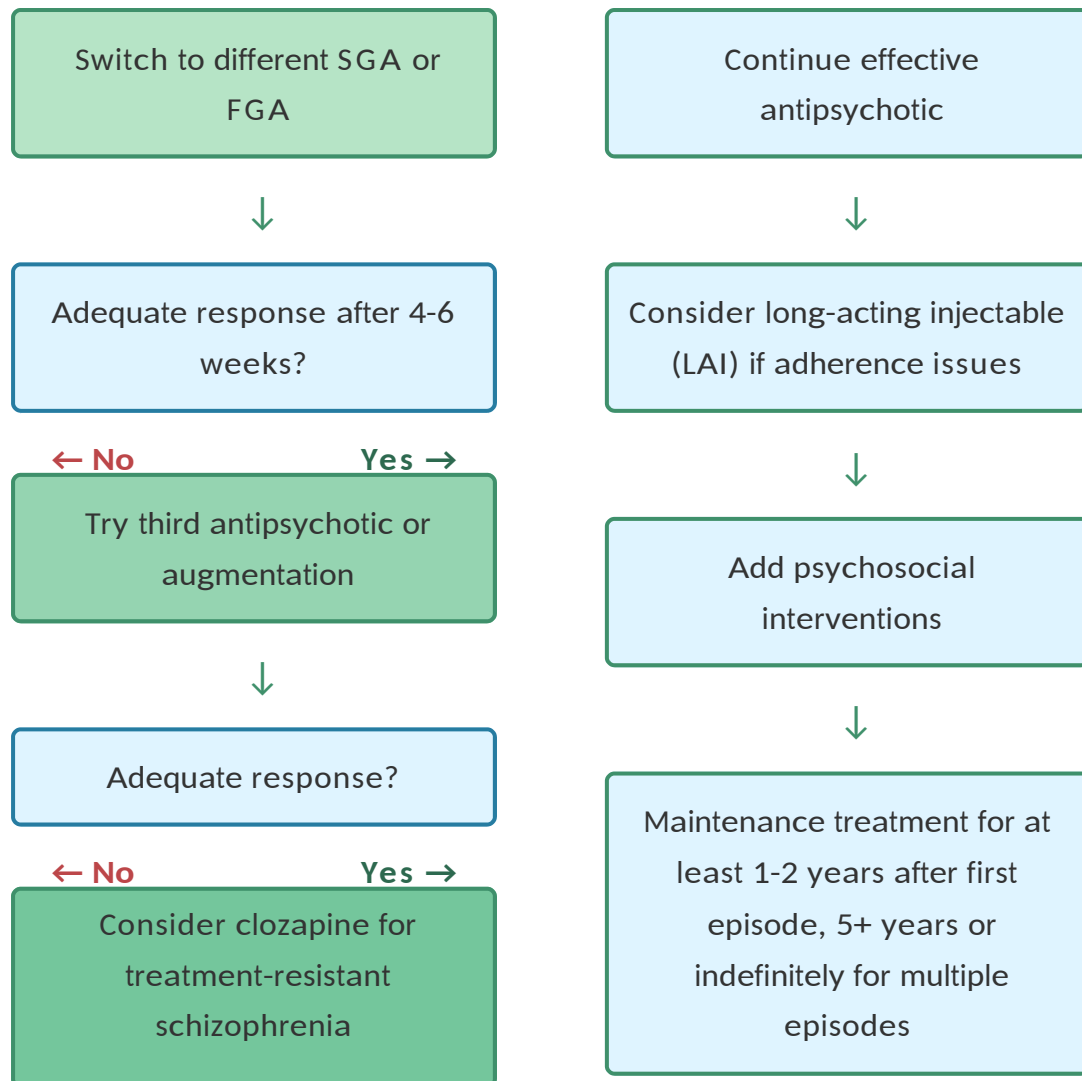
First-line: Oral SGA



Adequate response after 4-6 weeks?

← No

Yes →



### First-Line Treatment Options

#### Second-Generation Antipsychotics (SGAs):

- Risperidone
- Olanzapine
- Quetiapine
- Ziprasidone
- Aripiprazole
- Paliperidone
- Lurasidone
- Brexpiprazole
- Cariprazine

## Second-Line Treatment Options

### First-Generation Antipsychotics (FGAs):

- Haloperidol
- Perphenazine
- Fluphenazine
- Loxapine

**Alternative SGA not tried in first-line**

## Third-Line Treatment Options

### Augmentation strategies:

- Antipsychotic + mood stabilizer (valproate, lithium)
- Combination of two antipsychotics
- Add lamotrigine (for negative symptoms)

## Treatment-Resistant Schizophrenia Options

**Clozapine** (after failure of at least 2 adequate antipsychotic trials)

### Clozapine augmentation:

- Add another antipsychotic
- Add mood stabilizer
- Add ECT (for catatonia or persistent positive symptoms)
- Add lamotrigine or glycine (for negative symptoms)

## Long-Acting Injectable (LAI) Antipsychotics

### Consider for:

- History of non-adherence
- Patient preference
- Multiple relapses

### Options:

- Risperidone LAI (2 weeks)
- Paliperidone palmitate (1 month or 3 months)
- Aripiprazole monohydrate (1 month)
- Aripiprazole lauroxil (1-2 months)
- Olanzapine pamoate (2-4 weeks)
- Haloperidol decanoate (4 weeks)
- Fluphenazine decanoate (2-4 weeks)

### Medication Dosing Guidelines

Medication	Starting Dose	Therapeutic Dose Range	Maximum Dose
Risperidone	1-2 mg/day	2-6 mg/day	16 mg/day
Olanzapine	5-10 mg/day	10-20 mg/day	30 mg/day
Quetiapine	25-50 mg BID	300-750 mg/day	800 mg/day
Aripiprazole	5-10 mg/day	10-30 mg/day	30 mg/day
Ziprasidone	20 mg BID	40-80 mg BID	160 mg/day
Paliperidone	3-6 mg/day	3-12 mg/day	12 mg/day
Lurasidone	40 mg/day	40-80 mg/day	160 mg/day
Haloperidol	1-2 mg BID	5-20 mg/day	30 mg/day
Clozapine	12.5-25 mg/day	300-450 mg/day	900 mg/day

### Monitoring Requirements

#### All antipsychotics:

- Baseline: Weight, BMI, waist circumference, BP, fasting glucose, lipid panel, ECG (if cardiac risk)
- Weight: Monthly for 3 months, then quarterly
- Metabolic parameters: At 3 months, then annually
- AIMS for EPS: Every 6 months

### Clozapine-specific monitoring:

- ANC weekly for 6 months, then every 2 weeks for 6 months, then monthly
- Myocarditis monitoring: ECG and troponin at baseline and weekly for first month
- Consider clozapine levels if response inadequate or toxicity suspected

**Warning: Clozapine requires registration in a monitoring program due to risk of agranulocytosis. Olanzapine pamoate can cause post-injection delirium/sedation syndrome requiring 3-hour observation.**

**Antipsychotics increase mortality in elderly patients with dementia-related psychosis.**

***Clinical Pearl:** Negative symptoms and cognitive impairment often respond poorly to antipsychotics. Consider psychosocial interventions (cognitive remediation, social skills training) for these domains. SGAs may have advantages for negative symptoms compared to FGAs, but the difference is modest.*



## Generalized Anxiety Disorder (GAD) Treatment Algorithm

Initial Evaluation & Diagnosis of GAD



Assess Severity, Comorbidities, Previous Treatments



First-line: SSRI, SNRI, or Psychotherapy (CBT)



Adequate response after 4-6 weeks?

← No

Yes →

Switch to different SSRI/SNRI  
or add psychotherapy

Continue treatment for 12+  
months



Adequate response?

Assess for discontinuation

← No

Yes →

Try pregabalin, buspirone, or  
hydroxyzine

Taper medication slowly if  
discontinuing



Adequate response?

Monitor for relapse

← No

Yes →

Consider TCA, MAOI, or  
augmentation strategies

### First-Line Treatment Options

#### SSRIs:

- Escitalopram
- Paroxetine
- Sertraline

### **SNRIs:**

- Venlafaxine XR
- Duloxetine

### **Psychotherapy:**

- Cognitive Behavioral Therapy (CBT)
- Acceptance and Commitment Therapy (ACT)

## **Second-Line Treatment Options**

**Alternative SSRI/SNRI not tried in first-line**

**Pregabalin**

**Buspirone**

## **Third-Line Treatment Options**

**Hydroxyzine**

**Tricyclic Antidepressants (TCAs):**

- Imipramine
- Amitriptyline

**Quetiapine (low dose)**

## **Fourth-Line/Treatment-Resistant Options**

**Monoamine Oxidase Inhibitors (MAOIs):**

- Phenelzine

**Augmentation strategies:**

- Add pregabalin to SSRI/SNRI
- Add low-dose antipsychotic
- Add buspirone to SSRI/SNRI



## Short-term Adjunctive Treatment

### Benzodiazepines (short-term use only, 2-4 weeks):

- Alprazolam
- Clonazepam
- Diazepam
- Lorazepam

**Warning: Risk of dependence, tolerance, withdrawal, cognitive impairment. Not recommended for long-term use or in patients with history of substance use disorders.**

## Medication Dosing Guidelines

Medication	Starting Dose	Therapeutic Dose Range	Maximum Dose
Escitalopram	5-10 mg/day	10-20 mg/day	20 mg/day
Paroxetine	10-20 mg/day	20-50 mg/day	60 mg/day
Sertraline	25-50 mg/day	50-200 mg/day	200 mg/day
Venlafaxine XR	37.5-75 mg/day	75-225 mg/day	225 mg/day
Duloxetine	30 mg/day	60-120 mg/day	120 mg/day
Pregabalin	75 mg BID	300-600 mg/day	600 mg/day
Buspirone	5 mg BID	15-30 mg/day	60 mg/day
Hydroxyzine	25 mg TID-QID	50-100 mg TID-QID	400 mg/day

## Monitoring Requirements

### SSRIs/SNRIs:

- Assess for response and side effects at 2, 4, 8, and 12 weeks

- Monitor for activation, suicidality (especially in young adults)
- Check BP with SNRIs, especially venlafaxine

#### **Pregabalin:**

- Monitor for sedation, dizziness, weight gain
- Adjust dose in renal impairment

#### **Benzodiazepines:**

- Assess for sedation, cognitive impairment, dependence
- Limit duration of use
- Plan for gradual taper when discontinuing

**Clinical Pearl:** Response to pharmacotherapy for GAD may take 4-6 weeks for full effect. Combining medication with CBT often produces better outcomes than either treatment alone. Pregabalin has faster onset of action (1-2 weeks) compared to SSRIs/SNRIs.



## Obsessive-Compulsive Disorder (OCD) Treatment Algorithm

Initial Evaluation & Diagnosis of OCD

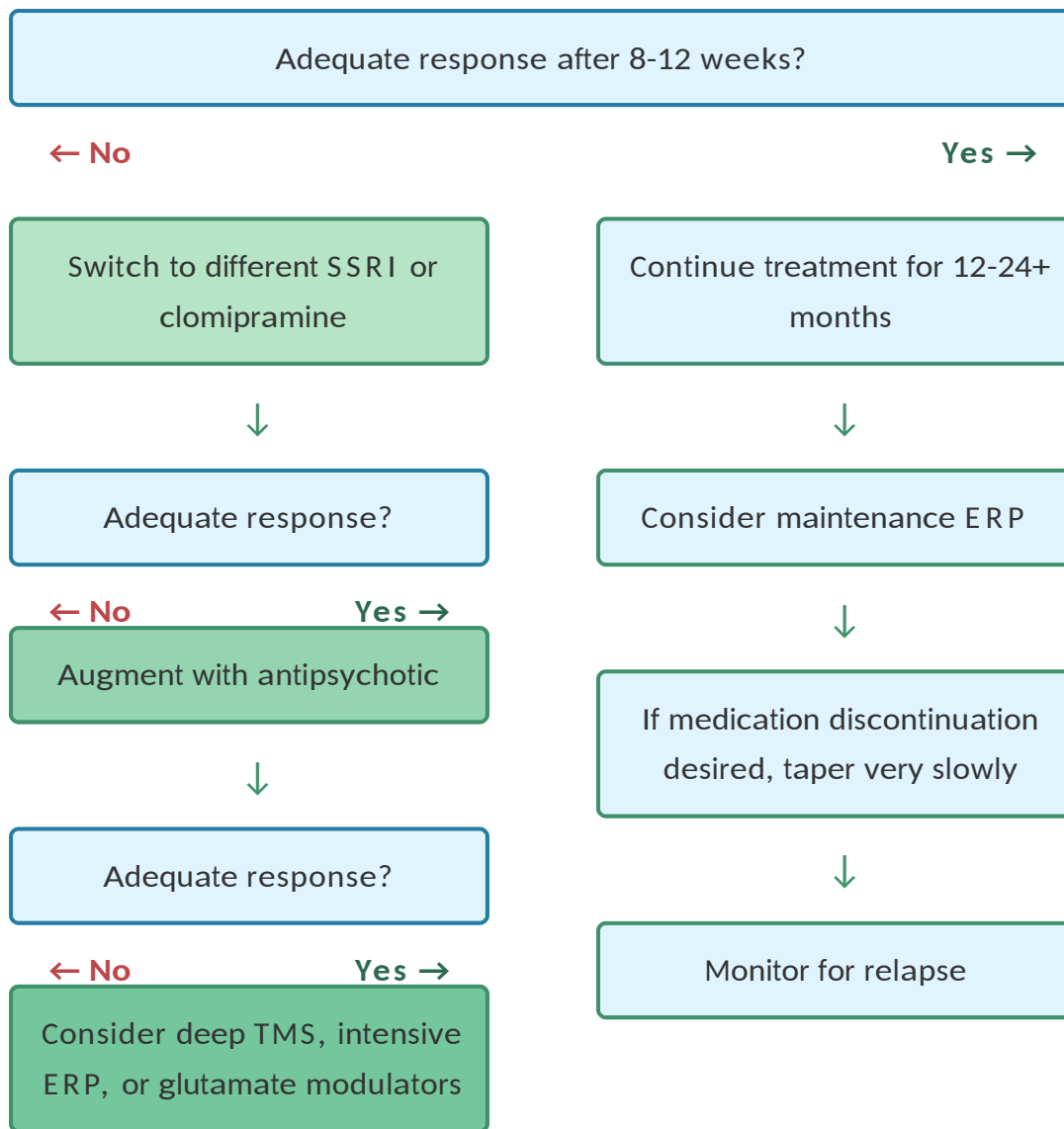


Assess Severity, Insight, Comorbidities



First-line: SSRI (higher doses) and/or ERP





### First-Line Treatment Options

#### SSRIs (higher doses than for depression):

- Fluoxetine
- Sertraline
- Paroxetine
- Fluvoxamine
- Escitalopram

#### Psychotherapy:

- Exposure and Response Prevention (ERP)

## Second-Line Treatment Options

**Alternative SSRI not tried in first-line**

**Clomipramine** (TCA with strong serotonergic effects)

## Third-Line/Augmentation Strategies

### Antipsychotic augmentation:

- Risperidone
- Aripiprazole
- Quetiapine
- Olanzapine

### Other augmentation strategies:

- Add clomipramine to SSRI (with caution)
- Add another SSRI (with caution)

## Treatment-Resistant OCD Options

### Glutamate modulators:

- Memantine
- N-acetylcysteine
- Riluzole

### Other options:

- Deep Transcranial Magnetic Stimulation (dTMS)
- Intensive residential treatment with ERP
- Gamma knife capsulotomy
- Deep brain stimulation

## Medication Dosing Guidelines

Medication	Starting Dose	Therapeutic Dose Range for OCD	Maximum Dose
------------	---------------	--------------------------------	--------------

Fluoxetine	20 mg/day	40-80 mg/day	120 mg/day
Sertraline	50 mg/day	100-200 mg/day	400 mg/day
Paroxetine	20 mg/day	40-60 mg/day	80 mg/day
Fluvoxamine	50 mg/day	200-300 mg/day	400 mg/day
Escitalopram	10 mg/day	20-40 mg/day	40 mg/day
Clomipramine	25 mg/day	150-250 mg/day	300 mg/day
Risperidone (augmentation)	0.5 mg/day	1-3 mg/day	4 mg/day
Aripiprazole (augmentation)	2.5 mg/day	5-15 mg/day	15 mg/day

## Monitoring Requirements

### SSRIs:

- Assess for response and side effects at 4, 8, and 12 weeks
- OCD often requires higher doses and longer trial duration (10-12 weeks)

### Clomipramine:

- Baseline: ECG (QTc interval)
- Follow-up ECG with dose increases
- Consider blood levels if response inadequate or toxicity suspected

### Antipsychotics:

- Baseline: Weight, BMI, waist circumference, BP, fasting glucose, lipid panel
- Weight: Monthly for 3 months, then quarterly
- Metabolic parameters: At 3 months, then annually
- AIMS for EPS: Every 6 months

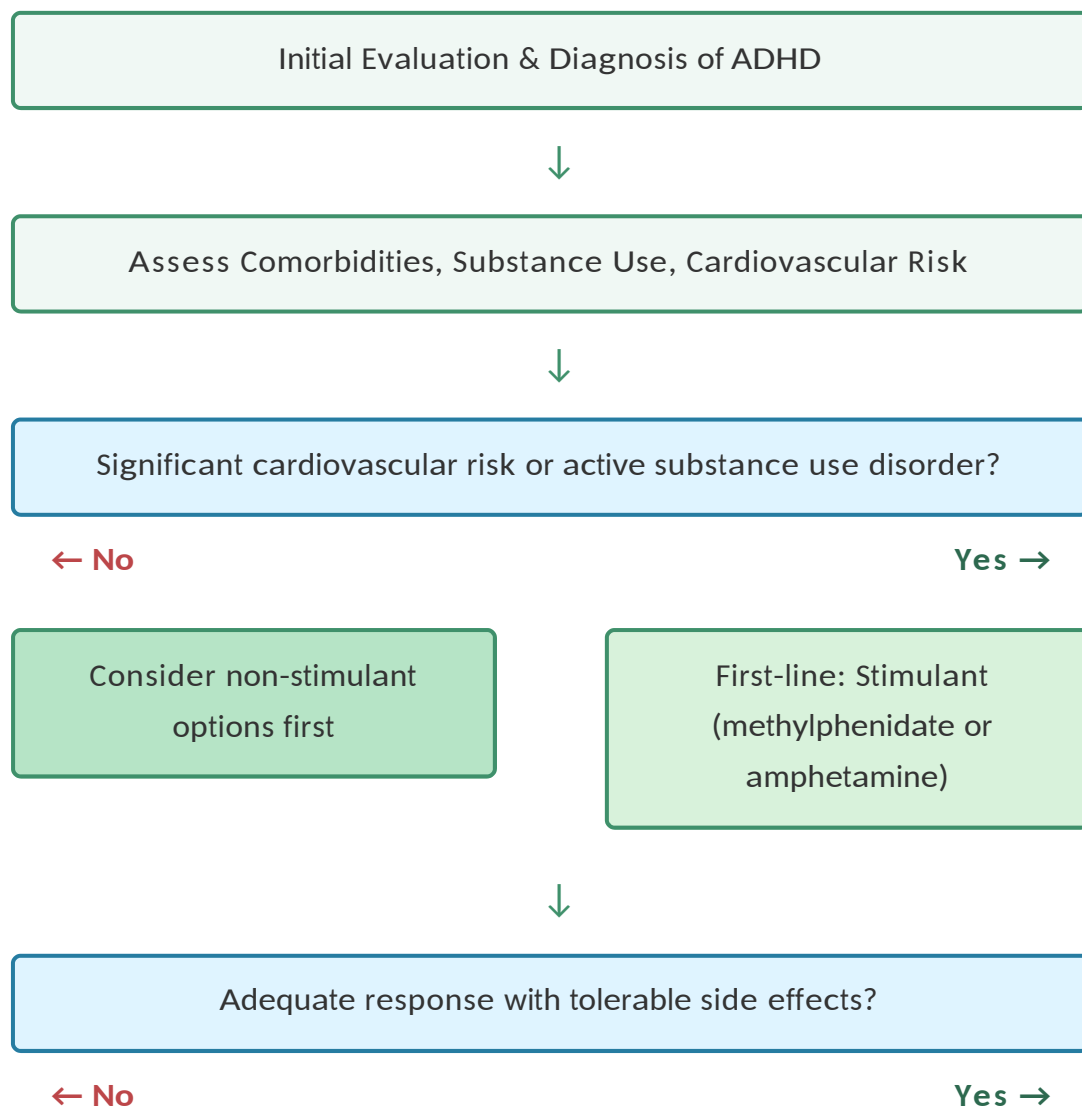
**Warning: Combining clomipramine with SSRIs can increase risk of serotonin syndrome. Abrupt discontinuation of OCD medications can**

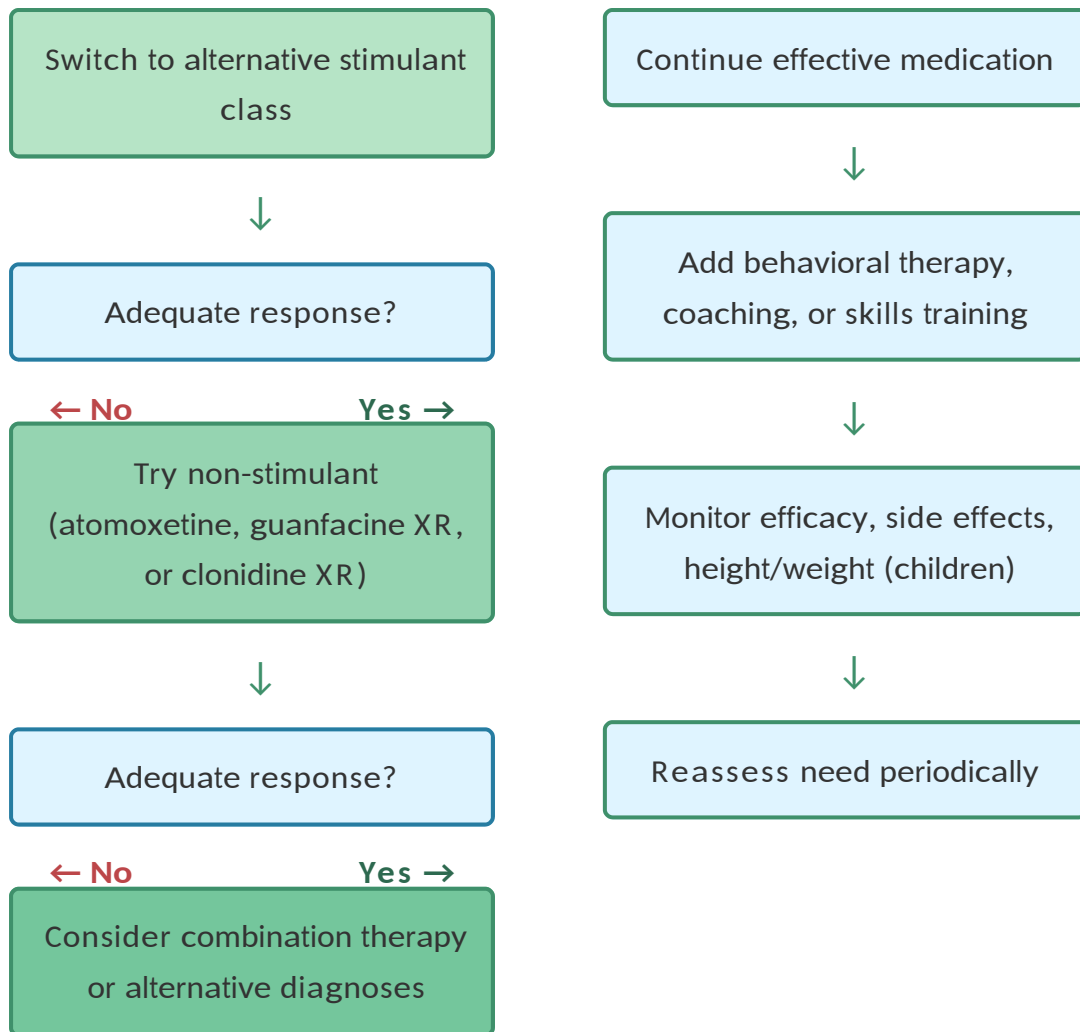
lead to high relapse rates and withdrawal symptoms. Taper very slowly when discontinuing.

**Clinical Pearl:** OCD typically requires higher SSRI doses than depression and longer trial duration (10-12 weeks). Combining medication with ERP produces better outcomes than either treatment alone. Poor insight OCD may respond better to antipsychotic augmentation.



## ADHD Treatment Algorithm





## First-Line Treatment Options

### Stimulants:

#### Methylphenidate-based:

- Methylphenidate IR, SR, ER
- Methylphenidate OROS (Concerta)
- Dexamethylphenidate IR, XR

#### Amphetamine-based:

- Mixed amphetamine salts IR, XR
- Lisdexamfetamine
- Dextroamphetamine IR, SR

## Second-Line Treatment Options

### Non-stimulants:

- Atomoxetine
- Guanfacine E R
- Clonidine E R

## Third-Line/Combination Treatment Options

### Combination therapy:

- Stimulant + guanfacine E R
- Stimulant + atomoxetine

### Other options:

- Bupropion
- Viloxazine
- Modafinil (off-label)

## Medication Dosing Guidelines

Medication	Starting Dose	Titration	Typical Effective Dose	Maximum Dose
Methylphenidate IR	5-10 mg BID-TID	↑ 5-10 mg weekly	20-60 mg/ day	60 mg/day (children) 100 mg/day (adults)
Methylphenidate E R (Concerta)	18 mg QAM	↑ 18 mg weekly	36-54 mg/ day	72 mg/day
Mixed amphetamine salts XR	5-10 mg QAM	↑ 5-10 mg weekly	15-30 mg/ day	30 mg/day (children) 60 mg/day (adults)



Lisdexamfetamine	30 mg QAM	↑ 10-20 mg weekly	50-70 mg/ day	70 mg/day
Atomoxetine	0.5 mg/ kg/day	↑ after 3 days to 1.2 mg/kg/day	1.2-1.4 mg/ kg/day	1.4 mg/kg/ day or 100 mg/day
Guanfacine ER	1 mg QD	↑ 1 mg weekly	2-4 mg/day	4 mg/day (children) 7 mg/day (adults)
Clonidine ER	0.1 mg QHS	↑ 0.1 mg weekly	0.2-0.4 mg/ day	0.4 mg/day

## Monitoring Requirements

### Stimulants:

- Baseline: BP, HR, height, weight
- Follow-up: BP, HR at each visit
- Height/weight: Every 3-6 months in children/adolescents
- Consider ECG if cardiac risk factors

### Atomoxetine:

- Baseline: BP, HR, LFTs if liver concerns
- Follow-up: BP, HR at each visit
- Monitor for suicidal ideation, especially early in treatment

### Guanfacine/Clonidine ER:

- Baseline: BP, HR, ECG
- Follow-up: BP, HR at each visit
- Monitor for sedation, dizziness, hypotension

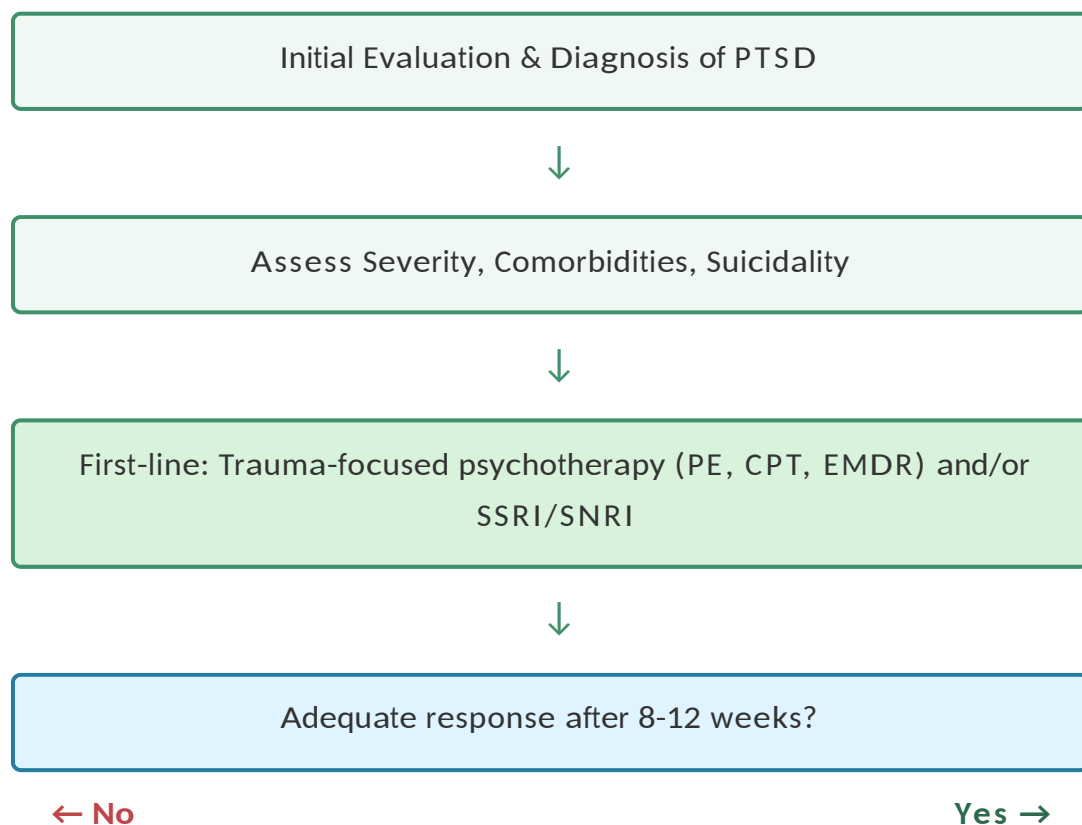
**Warning: Stimulants are controlled substances with potential for misuse. Avoid in patients with severe cardiovascular disease, uncontrolled**

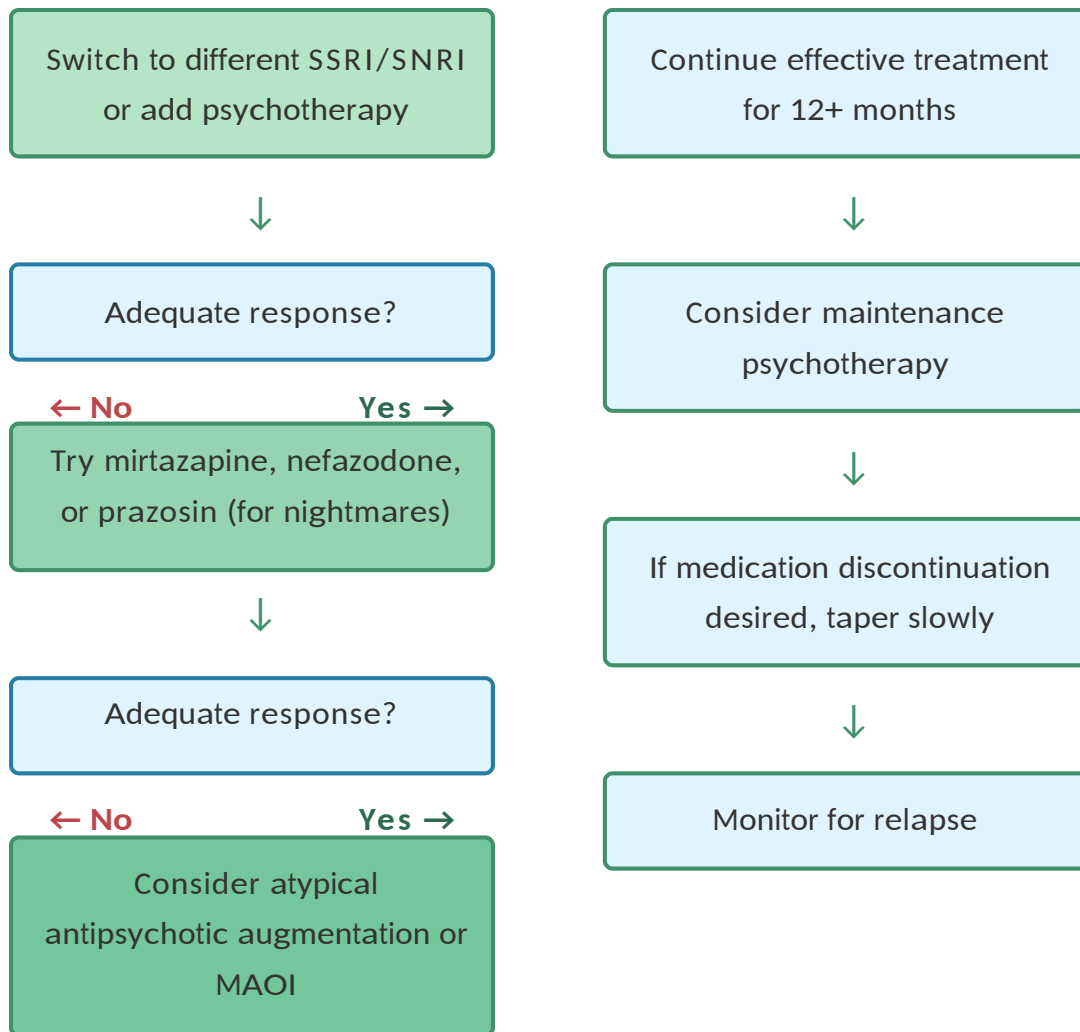
**hypertension, hyperthyroidism, or glaucoma. Stimulants may exacerbate tics, anxiety, or psychosis in vulnerable individuals.**

**Clinical Pearl:** About 70% of patients respond to the first stimulant trial. If one class of stimulant is ineffective or poorly tolerated, there is still a good chance of response to the other class. Combining medication with behavioral interventions typically produces better outcomes than medication alone.



## Post-Traumatic Stress Disorder (PTSD) Treatment Algorithm





### First-Line Treatment Options

#### Trauma-focused psychotherapy:

- Prolonged Exposure (PE)
- Cognitive Processing Therapy (CPT)
- Eye Movement Desensitization and Reprocessing (EMDR)

#### Medications:

- SSRIs: Sertraline, Paroxetine (FDA-approved for PTSD)
- Other SSRIs: Fluoxetine, Escitalopram
- SNRIs: Venlafaxine

### Second-Line Treatment Options

### **Alternative SSRI/SNRI not tried in first-line**

#### **Other antidepressants:**

- Mirtazapine
- Nefazodone

#### **For nightmares/sleep disturbance:**

- Prazosin

### **Third-Line/Augmentation Strategies**

#### **Antipsychotic augmentation:**

- Risperidone
- Quetiapine
- Olanzapine
- Aripiprazole

#### **Other options:**

- Mood stabilizers (lamotrigine, topiramate)
- Trazodone (for sleep)

### **Treatment-Resistant PTSD Options**

#### **MAOIs:**

- Phenelzine

#### **Other options:**

- Ketamine/Esketamine (investigational)
- MDMA-assisted psychotherapy (investigational)
- Stellate ganglion block
- Transcranial magnetic stimulation (TMS)

### **Medication Dosing Guidelines**

Medication	Starting Dose	Therapeutic Dose Range	Maximum Dose
Sertraline	25-50 mg/day	50-200 mg/day	200 mg/day
Paroxetine	10-20 mg/day	20-60 mg/day	60 mg/day
Fluoxetine	10-20 mg/day	20-60 mg/day	80 mg/day
Venlafaxine XR	37.5-75 mg/day	150-225 mg/day	375 mg/day
Mirtazapine	15 mg QHS	30-45 mg/day	45 mg/day
Prazosin	1 mg QHS	3-15 mg QHS	20 mg/day
Quetiapine (augmentation)	25-50 mg QHS	50-300 mg/day	300 mg/day
Risperidone (augmentation)	0.5 mg BID	1-3 mg/day	6 mg/day

### Monitoring Requirements

#### SSRIs/SNRIs:

- Assess for response and side effects at 2, 4, 8, and 12 weeks
- Monitor for activation, suicidality (especially in young adults)
- Check BP with SNRIs, especially venlafaxine

#### Prazosin:

- Monitor BP, especially with first dose (risk of first-dose hypotension)
- Start at low dose and titrate slowly

#### Antipsychotics:

- Baseline: Weight, BMI, waist circumference, BP, fasting glucose, lipid panel

- Weight: Monthly for 3 months, then quarterly
- Metabolic parameters: At 3 months, then annually
- AIMS for EPS: Every 6 months

**Warning: Benzodiazepines are generally not recommended for PTSD as they may interfere with fear extinction and trauma processing. They may worsen outcomes in the long term and carry risks of dependence and withdrawal.**

**Clinical Pearl:** Trauma-focused psychotherapies have the strongest evidence base for PTSD treatment. Combining medication with psychotherapy may be more effective than either alone, especially for complex or severe PTSD. Prazosin is particularly effective for trauma-related nightmares but less so for daytime symptoms.

### **General Clinical Pearls for Psychiatric Treatment:**

- Start low, go slow with medication dosing, especially in elderly, medically complex patients, or those sensitive to side effects
- Adequate trial duration is essential before concluding treatment failure (4-6 weeks for most conditions, 10-12 weeks for OCD)
- Combining pharmacotherapy with appropriate psychotherapy typically produces better outcomes than either treatment alone
- Consider patient preferences, prior treatment response, side effect profiles, and comorbidities when selecting treatments
- Monitor for suicidality with all psychotropic medications, especially during initiation and dose changes
- Treatment adherence is critical; address barriers and consider long-acting formulations when appropriate
- Taper medications slowly when discontinuing to minimize withdrawal symptoms and relapse risk

### **References**

- Bains, N., & Abdijadid, S. (2023, April 10). *Major depressive disorder*. PubMed; StatPearls Publishing.  
<https://www.ncbi.nlm.nih.gov/books/NBK559078/>
- Beaulieu, A. M., Tabasky, E., & Osser, D. N. (2019). The psychopharmacology algorithm project at the Harvard South Shore Program: An algorithm for adults with obsessive-compulsive disorder. *Psychiatry Research*, 281, 112583. <https://doi.org/10.1016/j.psychres.2019.112583>
- DeGeorge, K. C., Grover, M., & Streeter, G. S. (2022). Generalized anxiety disorder and panic disorder in adults. *American Family Physician*, 106(2), 157–164.  
<https://www.aafp.org/pubs/afp/issues/2022/0800/generalized-anxiety-disorder-panic-disorder.html>
- Eom, T.-H., & Kim, Y.-H. (2023). Clinical Practice Guidelines for Attention-deficit/Hyperactivity Disorder: Recent Updates. *Clinical and Experimental Pediatrics (Online)*, 67(1).  
<https://doi.org/10.3345/cep.2021.01466>
- H, C. (2011, November 16). *Pharmacotherapy of Major Depressive Disorder*. Uspharmacist.com.  
<https://www.uspharmacist.com/article/pharmacotherapy-of-major-depressive-disorder>
- Holtzheimer, P., & Montañó, M. (2014). *Clinician’s Guide to Medications for PTSD - PTSD: National Center for PTSD*. Va.gov.  
[https://www.ptsd.va.gov/professional/treat/txessentials/clinician\\_guide\\_meds.asp](https://www.ptsd.va.gov/professional/treat/txessentials/clinician_guide_meds.asp)
- Implementing the Key Action Statements: An Algorithm and Explanation for Process of Care for the Evaluation, Diagnosis, Treatment, and Monitoring of ADHD in Children and Adolescents*. (n.d.).  
[https://chsciowa.org/sites/chsciowa.org/files/resource/files/3\\_-\\_adhd\\_algorithm\\_supplement.pdf](https://chsciowa.org/sites/chsciowa.org/files/resource/files/3_-_adhd_algorithm_supplement.pdf)
- Jain, A., & Mitra, P. (2023, February 20). *Bipolar Disorder*. PubMed; StatPearls Publishing.  
<https://www.ncbi.nlm.nih.gov/books/NBK558998/>
- Kimmel, R. (2018). *UW PACC Psychiatry and Addictions Case Conference MEDICATION ALGORITHM FOR ANXIETY DISORDERS*.  
[https://ictp.uw.edu/sites/default/files/UWPACC\\_2018\\_10\\_18\\_Anxiety\\_An\\_Algorithm\\_for\\_medication\\_trials\\_Ryan\\_Kimmel\\_MD.pdf](https://ictp.uw.edu/sites/default/files/UWPACC_2018_10_18_Anxiety_An_Algorithm_for_medication_trials_Ryan_Kimmel_MD.pdf)

- Kishi, T., Ikuta, T., Matsuda, Y., Sakuma, K., Okuya, M., Nomura, I., Hatano, M., & Iwata, N. (2021). Pharmacological treatment for bipolar mania: A systematic review and network meta-analysis of double-blind randomized controlled trials. *Molecular Psychiatry*, 27(2).  
<https://doi.org/10.1038/s41380-021-01334-4>
- Kovich, H., Kim, W., & Quaste, A. M. (2023). Pharmacologic Treatment of Depression. *American Family Physician*, 107(2), 173–181.  
<https://www.aafp.org/pubs/afp/issues/2023/0200/pharmacologic-treatment-of-depression.html>
- Martin, A., Naunton, M., Kosari, S., Peterson, G., Thomas, J., & Christenson, J. K. (2021). Treatment guidelines for PTSD: A systematic review. *Journal of Clinical Medicine*, 10(18).  
<https://doi.org/10.3390/jcm10184175>
- Nazarova, V. A., Sokolov, A. V., Chubarev, V. N., Tarasov, V. V., & Schiöth, H. B. (2022). Treatment of ADHD: Drugs, Psychological therapies, devices, Complementary and Alternative Methods as Well as the Trends in Clinical Trials. *Frontiers in Pharmacology*, 13(13), 1–19.  
<https://doi.org/10.3389/fphar.2022.1066988>
- Patel, K. R., Cherian, J., Gohil, K., & Atkinson, D. (2014). Schizophrenia: Overview and Treatment Options. *Pharmacy and Therapeutics*, 39(9), 638.  
<https://pmc.ncbi.nlm.nih.gov/articles/PMC4159061/>
- Paxos, C. (2022). Moving beyond first-line treatment options for OCD. *Mental Health Clinician*, 12(5), 300–308. <https://doi.org/10.9740/mhc.2022.10.300>
- Rybakowski, J. K. (2023). Application of Antipsychotic Drugs in Mood Disorders. *Brain Sciences*, 13(3), 414. <https://doi.org/10.3390/brainsci13030414>
- Sartor, Z., Kelley, L., & Laschober, R. (2023). Posttraumatic Stress Disorder: Evaluation and Treatment. *American Family Physician*, 107(3), 273–281.  
<https://www.aafp.org/pubs/afp/issues/2023/0300/posttraumatic-stress-disorder.html>
- Shah, N., Grover, S., & Rao, Gp. (2020). Clinical practice guidelines for management of bipolar disorder. *Indian Journal of Psychiatry*, 59(5), 51. <https://doi.org/10.4103/0019-5545.196974>